

fMRI and MRI at NIH

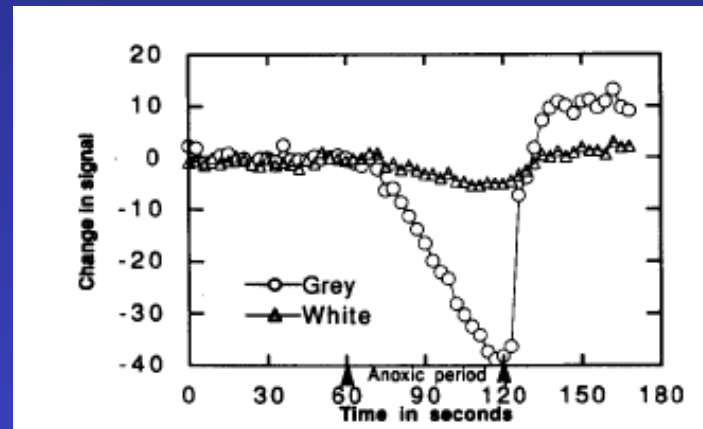
Sean Marrett / FMRI / NIMH

MRI and Brain MRI at NIH

1. Development of MRI at NIH
 - In-Vivo NMR Center (established 1986ish)
 - 2 Bruker animal magnets, GE Signa 1.5T
 - 4 Tesla with head gradient (NHLBI 1989ish)
2. Early BOLD studies in animals
3. Initial human functional studies (1993ish)
4. Key developments from NIH and MRI researchers now at NIH
 - DTI
 - Magnetization Transfer
 - Population-based longitudinal studies of brain development
 - Imaging genomics
 - FMRI
 - Perfusion imaging (Arterial Spin Labeling)
 - Decoding/Multivoxel Pattern Analysis
 - High resolution anatomical imaging
 - Real-time FMRI / Analysis Software
 - High-field imaging (4 Tesla, 7 Tesla)

Early BOLD MR Measurements in Animals

Turner R, Le Bihan D, Moonen CT, Despres D, Frank J “Echo-planar time course MRI of cat brain oxygenation changes” Magn Reson Med. 1991 Nov;22(1):159-66



Early FMRI in Humans

Functional Mapping of the Human Visual Cortex at 4 and 1.5 Tesla Using Deoxygenation Contrast EPI

R. Turner, P. Jezzard, H. Wen, K. K. Kwong, D. Le Bihan, T. Zeffiro, R. S. Balaban

The effects of photic stimulation on the visual cortex of human brain were studied by means of gradient-echo echo-planar imaging (EPI). Whole-body 4 and 1.5 T MRI systems, equipped with a small z axis head gradient coil, gave an image intensity of up to 28% at 4 T, an increase observed in primary visual cortex, a decrease of blood oxygenation in regions of activity. The larger effects at 4 T are of importance due to the susceptibility differences between deoxygenated and oxygenated blood at high

field strength. The results show that the changes in blood flow are greater than in oxygen utilization during somatosensory stimulation. Similar results were reported in cat brain during electrical stimulation by Lübbers and Leni-

Functional Mapping

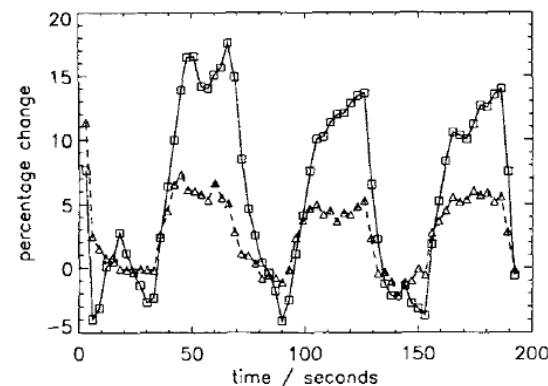


FIG. 2. Plot of fractional change in 4 T (squares) and 1.5 T (triangles) EPI image intensity versus time in the eight-voxel regions of interest in the visual cortex shown in Fig. 1, for a volunteer experiencing alternate 30-s periods of rest and photic stimulation. Details of acquisition for the 4 and 1.5 T data are described in the caption for Fig. 1.

agnetic fields the effect of susceptibility changes due to photic stimulation using our 4 T whole-body MR system. In comparison, EPI experiments at 1.5 T with the same subject

Invention of DTI

Biophysical Journal Volume 66 January 1994 259–267

259

MR Diffusion *Tensor* Spectroscopy and Imaging

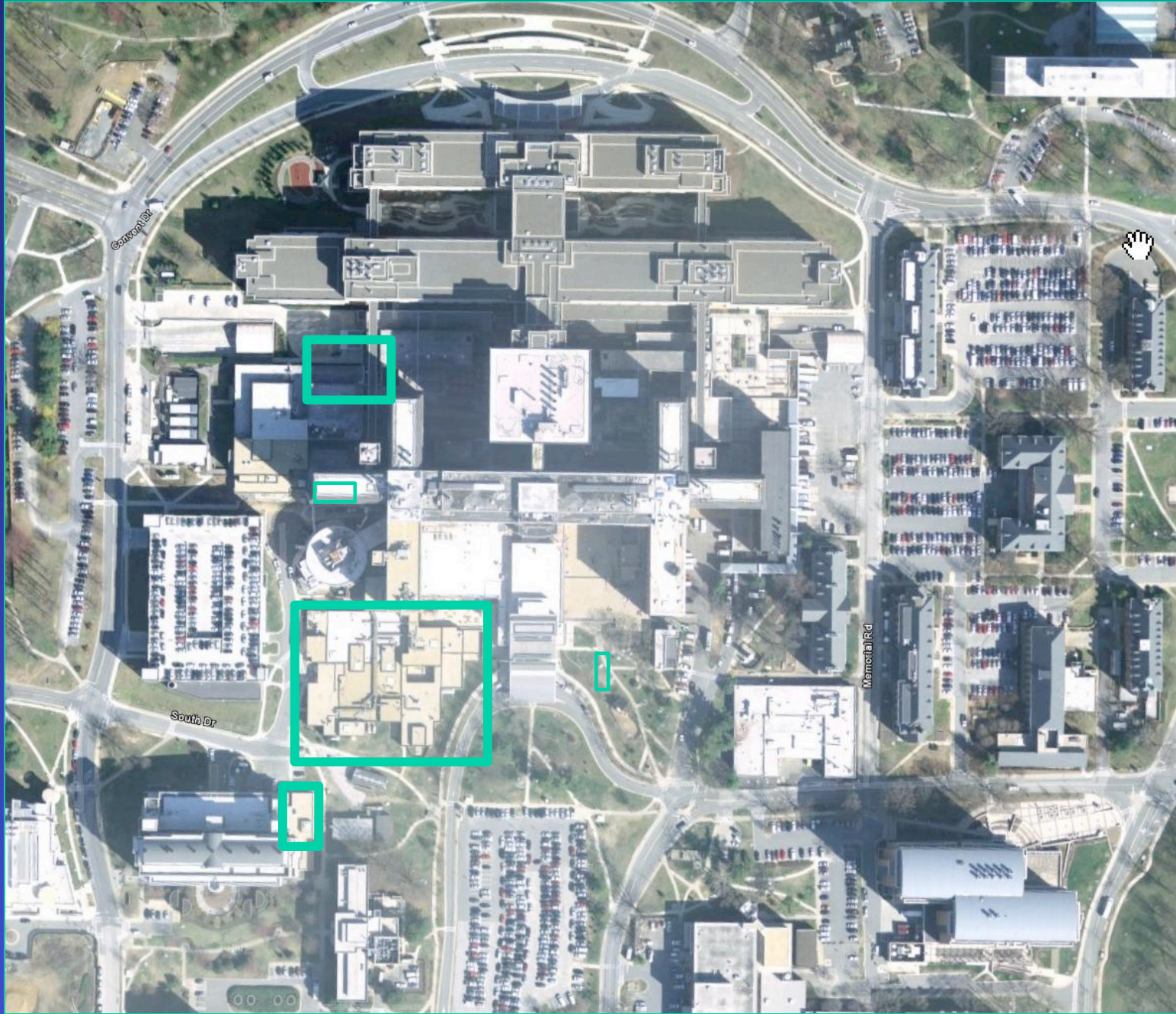
Peter J. Basser,* James Mattiello,* and Denis LeBihan†

*Biomedical Engineering and Instrumentation Program, National Center for Research Resources, and †Diagnostic Radiology Department, The Warren G. Magnuson Clinical Center, National Institutes of Health, Bethesda, Maryland 20892 USA

ABSTRACT This paper describes a new NMR imaging modality—MR diffusion *tensor* imaging. It consists of estimating an effective diffusion tensor, \mathbf{D}_{eff} , within a voxel, and then displaying useful quantities derived from it. We show how the phenomenon of anisotropic diffusion of water (or metabolites) in anisotropic tissues, measured noninvasively by these NMR methods, is exploited to determine fiber tract orientation and mean particle displacements. Once \mathbf{D}_{eff} is estimated from a series of NMR pulsed-gradient, spin-echo experiments, a tissue's three orthotropic axes can be determined. They coincide with the eigenvectors of \mathbf{D}_{eff} , while the effective diffusivities along these orthotropic directions are the eigenvalues of \mathbf{D}_{eff} . Diffusion ellipsoids, constructed in each voxel from \mathbf{D}_{eff} , depict both these orthotropic axes and the mean diffusion distances in these directions. Moreover, the three scalar invariants of \mathbf{D}_{eff} , which are independent of the tissue's orientation in the laboratory frame of reference, reveal useful information about molecular mobility reflective of local microstructure and anatomy. Inherently, tensors (like \mathbf{D}_{eff}) describing transport processes in anisotropic media contain new information *within a macroscopic voxel* that scalars (such as the apparent diffusivity, proton density, T_1 , and T_2) do not.

Where is MRI At NIH - Human

1. NIH MRI Research Facility (NMRF)
 - 3T GE HDx
2. FMRIF (NIMH & NINDS)
 - 3 x 3T GE HDx
 - 1 x 1.5T GE HDx
 - (in preparation - 2010) 7T Siemens/Magnex
 - (in preparation - 2011) 3T Siemens (Skyra)
3. NINDS/NIMH
 - 7T GE HD
 - (in preparation – 2011) 11.7T Siemens/Magnex
4. NINDS (Stroke)
 - Suburban Hospital
 - Washington Hospital
5. Clinical Center (Radiology & Imaging Sciences, TBI)
 - 3T & 1.5T Philips & 3T Siemens
6. NHLBI (Cardiac Energetics, Cardiovascular Intervention)
 - Multiple 1.5T Siemens Scanners
7. NCI/CCR – Molecular Imaging Program (also at Ft. Dietrich)



NIH NMR Facility



Where is fMRI at NIH - Human

1. NIH MRI Research Facility (NMRF)
 - 3T GE HDx
2. FMRI (NIMH & NINDS)
 - 3 x 3T GE HDx
 - 1 x 1.5T GE HDx
 - (in preparation - 2010) 7T Siemens/Magnex
 - (in preparation - 2011) 3T Siemens (Skyra)
3. NINDS/NIMH
 - 7T GE HD
 - (in preparation – 2011) 11.7T Siemens/Magnex

In-Vivo NMR Center Magnets



Where is fMRI at NIH (dedicated animal)

1. NIH MRI Research Facility (NMRF) & Mouse Imaging Facility
 - Multiple small animal magnets (4.7T, 7T Bruker)
2. Neurophysiology Imaging Facility (NIMH/NINDS/NEI)
 - dedicated 4.7T vertical bore primate
3. LFMI/NINDS
 - 7T Bruker (marmoset)
 - 11.7T Bruker (rodent, small animal)
4. NCI-Frederick Small Animal Imaging Program
 - 3T Philips

fMRI research + resources at the NIH

Studies focusing on:

- Normal Brain Function
- Mental disorders
 - Schizophrenia
 - Mood disorders
 - ADHD
 - Autism
 - Williams syndrome
 - Psychosis

Within FMRIIF 7 human MRI scanners (1 - 1.5T, 5 - 3T, 1 - 7T)

MEG, EEG

3 dedicated animal MRI scanners

Micro CT, Ultrasound, Bioluminescence

Ongoing fMRI Studies

- Epilepsy
- Visual processing
- Mood disorders
- Learning
- Schizophrenia/Imaging Genetics
- Plasticity/Recovery
- Motor Function
- Auditory processing
- Attention
- Language
- Speech
- Stroke
- Social Interaction
- Development
- Decision making

Users

NIMH:

Peter Bandettini, Ph.D.
Chris Baker, Ph.D.
Karen Berman, M.D.
James Blair, Ph.D.
Jay Giedd, M.D.
Christian Grillon, Ph.D.
Wayne Drevets, M.D.
Ellen Liebenluft, M.D.
Alex Martin, Ph.D.
Husseini Manji, M.D.
Andreas Meyer-Lindenberg, M.D.
Mort Mishkin, Ph.D.
Elizabeth Murray, Ph.D.
Daniel Pine, M.D.
Judith Rapaport, M.D.
Jun Shen, Ph.D.
Susan Swedo, M.D.
Leslie Ungerleider, Ph.D.
Daniel Weinberger, M.D.

NINDS:

Leonardo Cohen, M.D.
Jeff Duyn, Ph.D.
Jordan Grafman, Ph.D.
Mark Hallet, Ph.D.
John Hallenbeck, M.D.
Alan Koretsky, Ph.D.
Edward Oldfield, M.D.
Danny Reich, M.D., Ph.D.
William Theodore, M.D.

NIAAA:

Daniel Hommer, M.D.

NICHD:

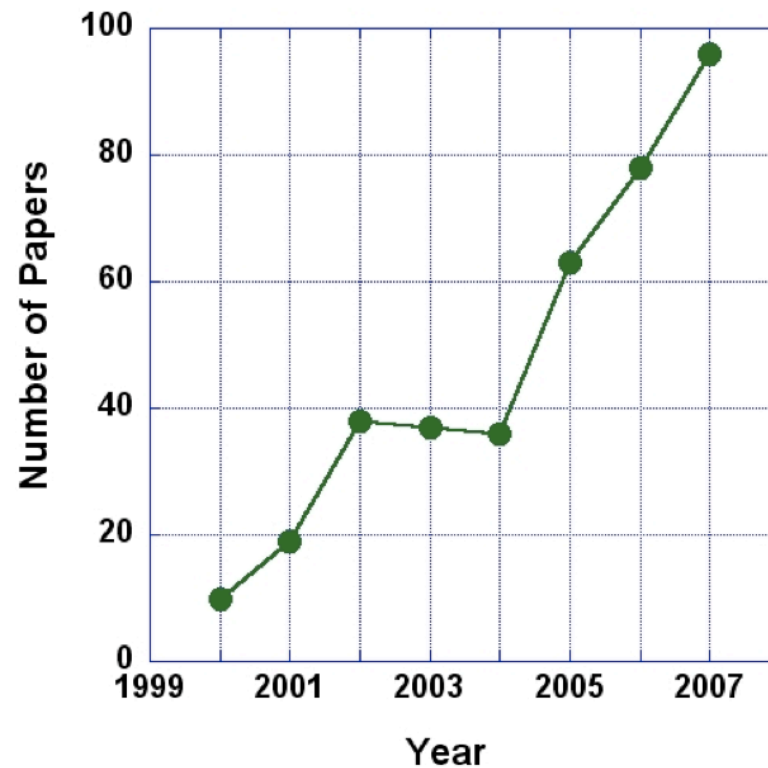
Peter Basser, Ph.D.
Allen Braun, M.D.

NCI:

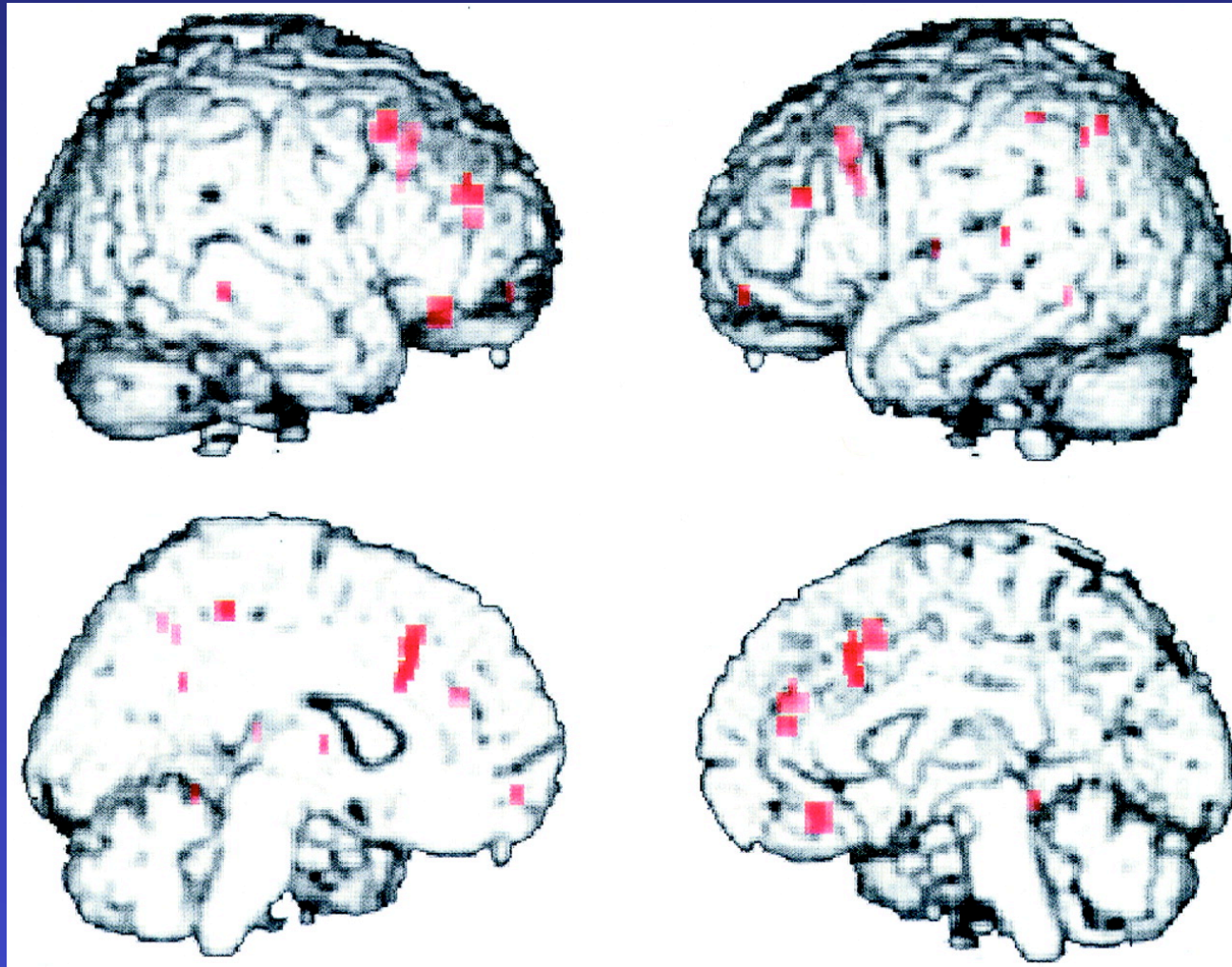
Kathy Warren, M.D.

Papers from FMRI

Number of Papers Produced per year by
Researchers using the Functional MRI Facility



Effect of COMT genotype on fMRI activation during the two-back working memory task.



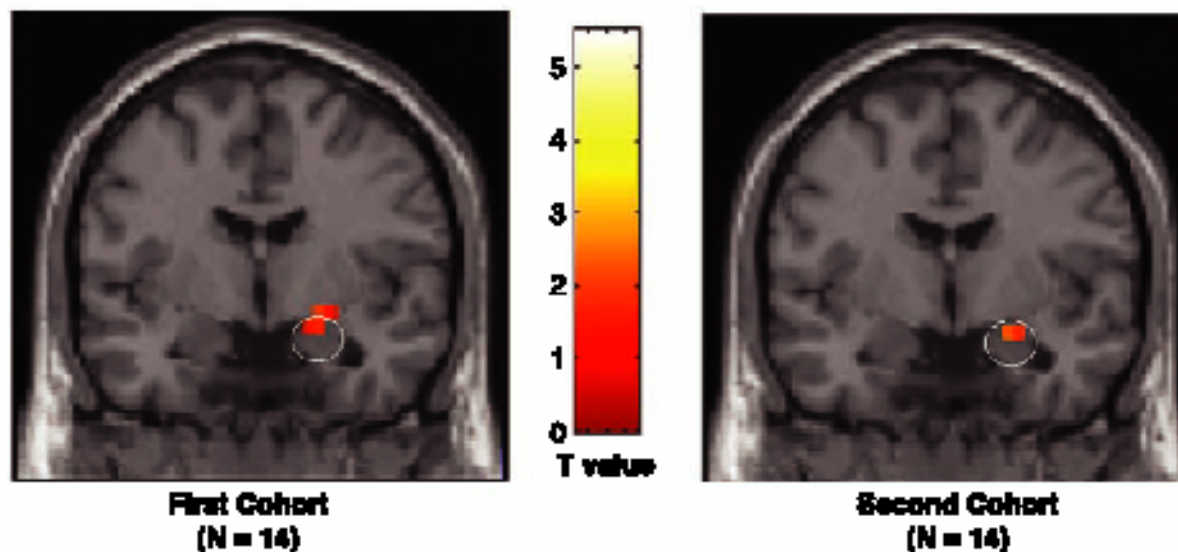
Egan M F et al. PNAS 2001;98:6917-6922

Comparison of two groups of *normal* individuals with differences in the Serotonin Transporter Gene

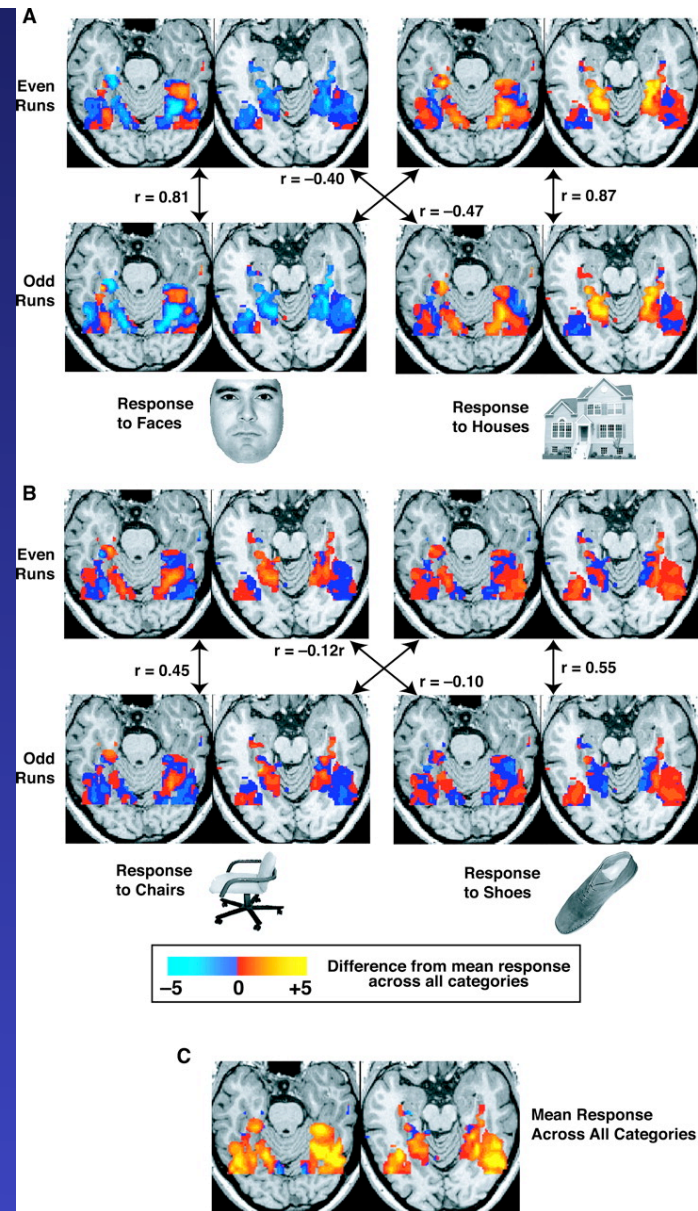
Serotonin Transporter Genetic Variation and the Response of the Human Amygdala

Ahmad R. Hariri,¹ Venkata S. Mattay,¹ Alessandro Tessitore,¹
Bhaskar Kolachana,¹ Francesco Fera,¹ David Goldman,²
Michael F. Egan,¹ Daniel R. Weinberger^{1*}

Amygdala Response: 2 Group > 1 Group



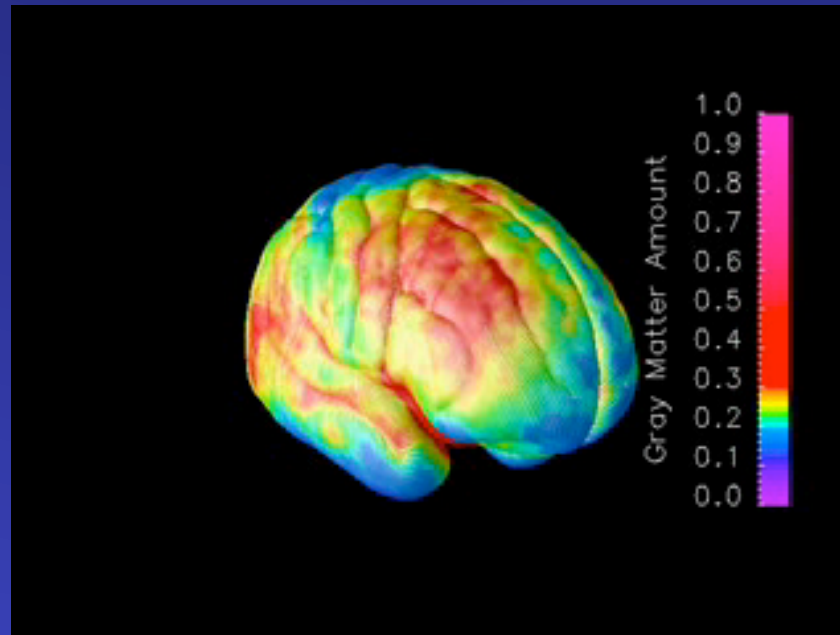
Decoding & Multivoxel Pattern Analysis



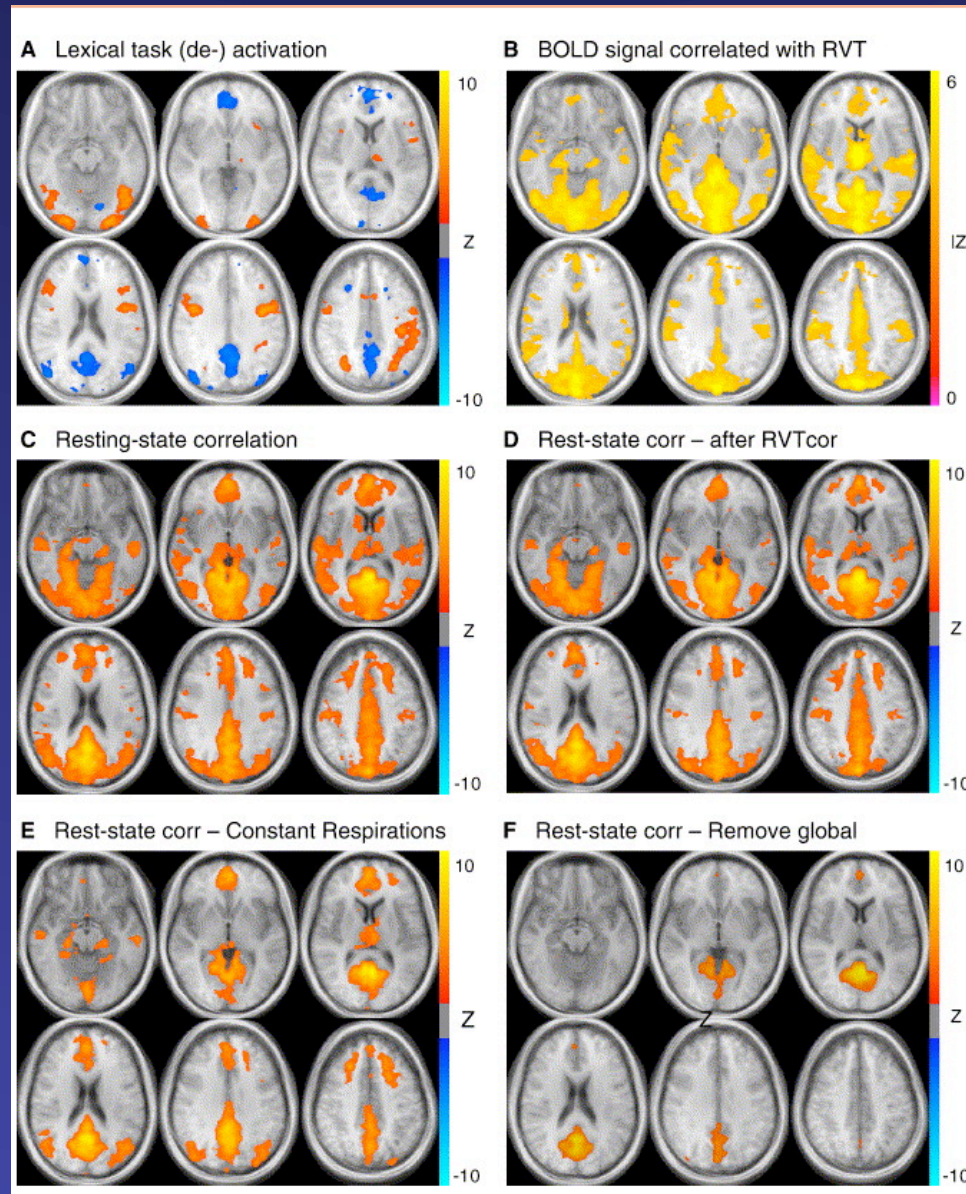
J. V. Haxby et al., Science 293, 2425 -2430 (2001)

Published by AAAS

Dynamic mapping of human cortical development during childhood through early adulthood (Gogtay et al. PNAS 2004)

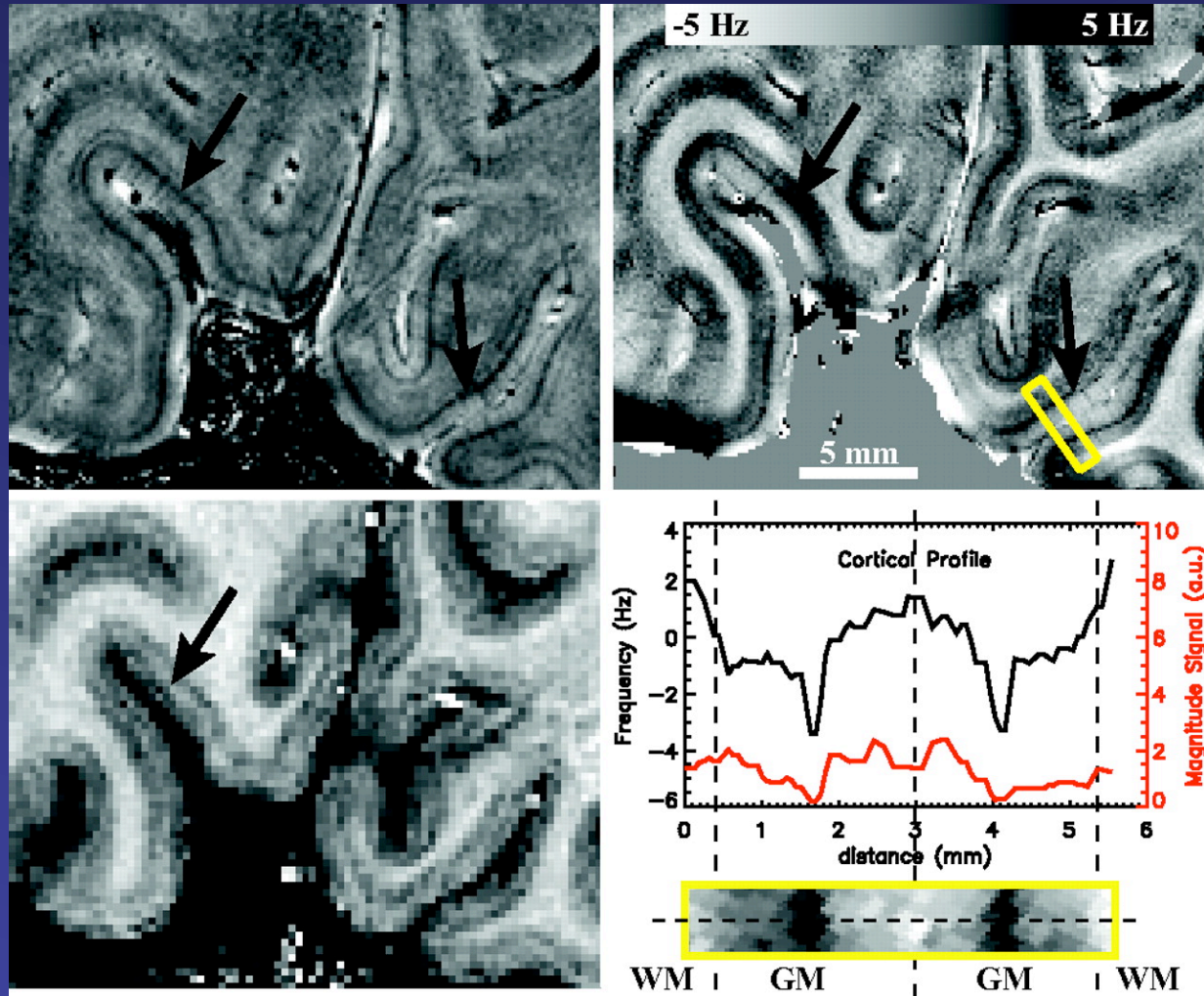


Separating respiratory-variation-related fluctuations from neuronal-activity-related fluctuations in fMRI



Birn et al, 2006

Intracortical contrast in the primary visual cortex.



Duyn J H et al. PNAS 2007;104:11796-11801

Functional MRI Facility



Souheil Inati staff scientist
Wenming Luh staff scientist
Sean Marrett staff scientist
Vinai Roopchansingh staff scientist
Adam Thomas system admin

Ellen Condon technologist
Janet Ebron technologist
Kenny Kan technologist
Marcela Montequin technologist
Sandra Moore technologist
Paula Rowser technologist
Debora Tkaczyk technologist
Jennifer Yeager technologist

Dorian Van Tassell admin lab manager

<http://fmrif.nimh.nih.gov/>

<http://fmrif.nih.gov/>



fmrif.nih.gov

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site navigation

public resources:

- About
- Research
- Investigators
- Staff
- Education
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- Directions
- NMRF Map
- fMRI Discussion

internal resources




- Help!
- scanner docs
- tools/software
- forums
- mailing lists
- faqs
- scheduling
- tech schedules

Welcome to the fMRI Facility at NIH

by Root — last modified 2005-11-03 11:53

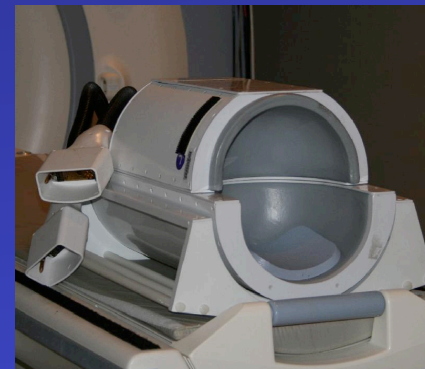
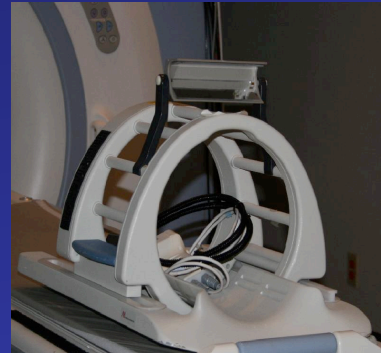
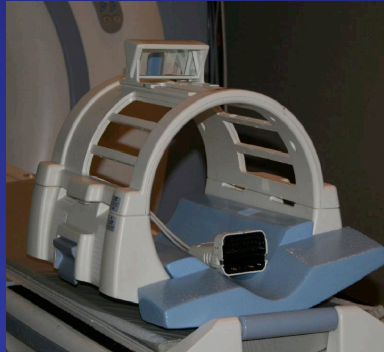


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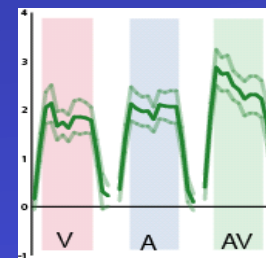
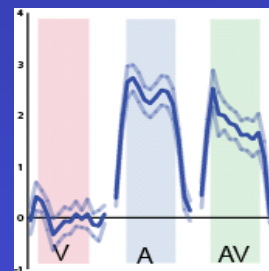
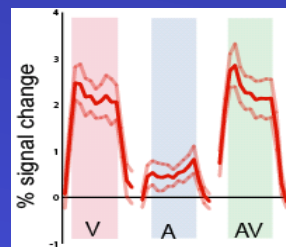
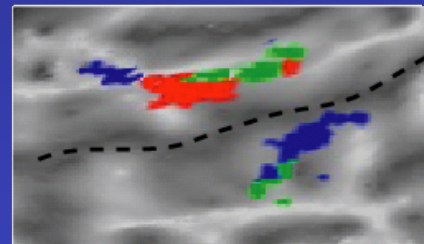
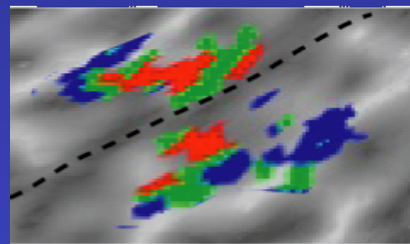
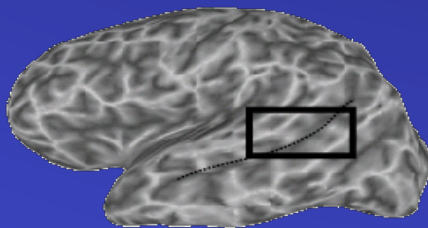
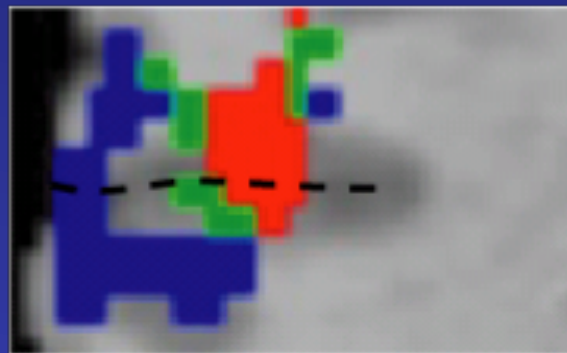
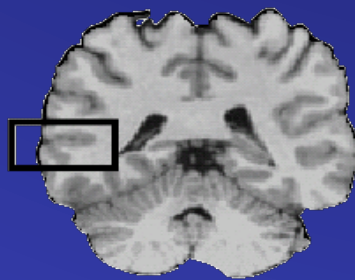
Basic Coil Options



Using Parallel Imaging to Increase Spatial Resolution of fMRI

Unravelling Multi-sensory integration

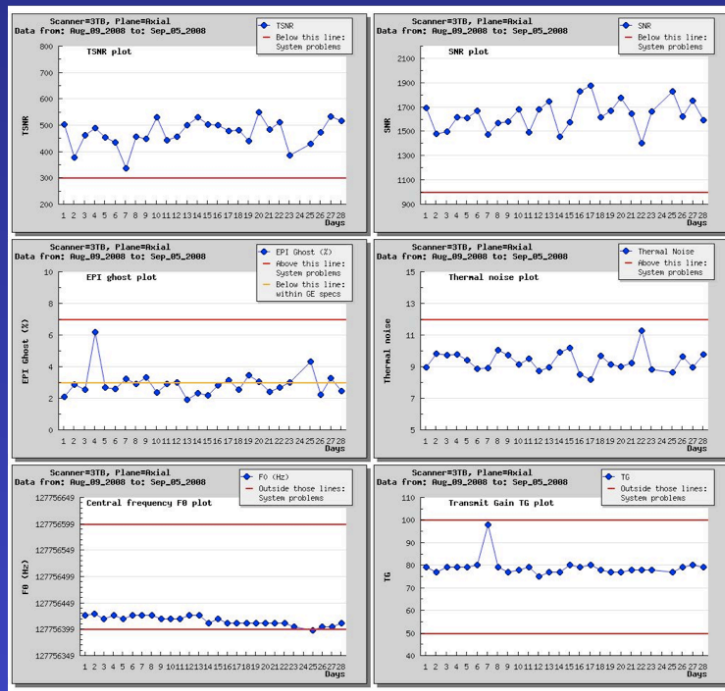
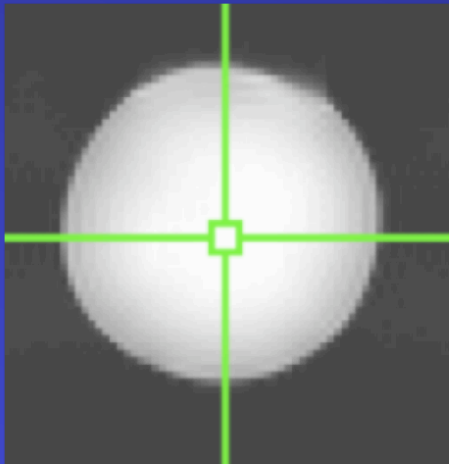
M.S. Beauchamp et al.,



Data and Q/A

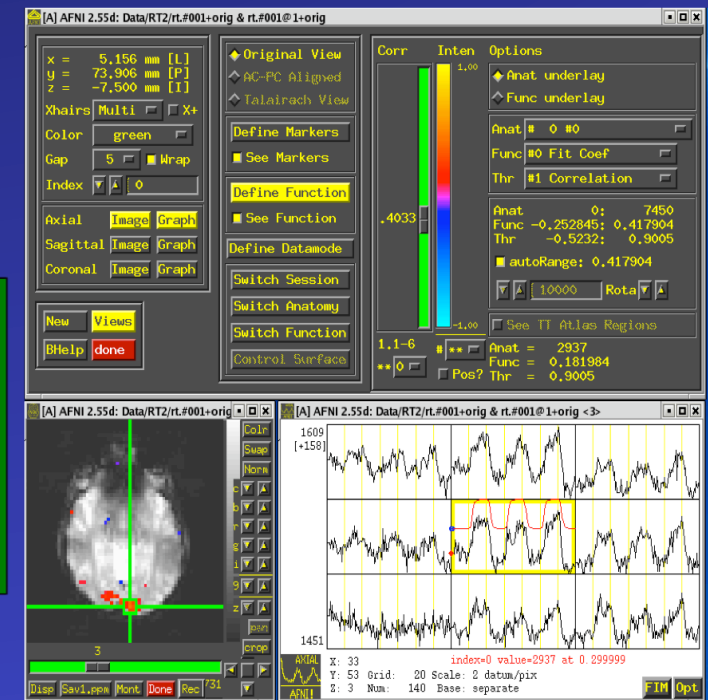
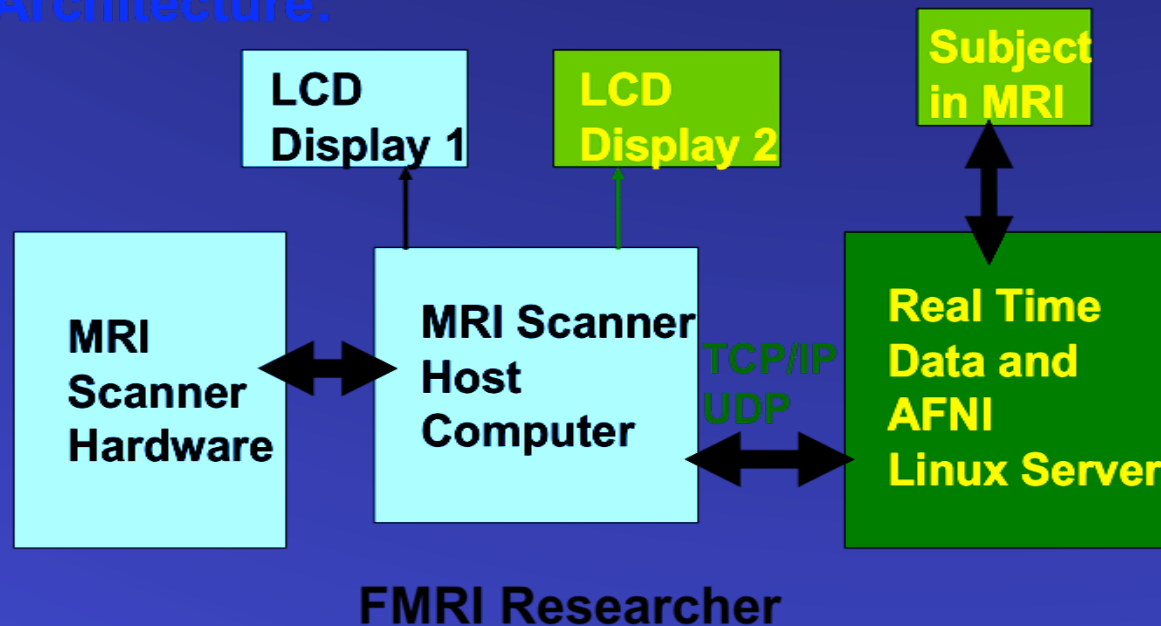
QA web interface

Quality Assurance (QA) Phantom



Real-time AFNI

Architecture:



Subject Interface



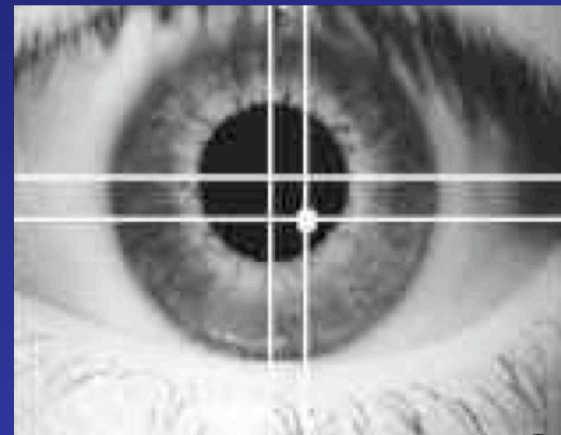
Avotec video



Rear Projection



Nordic Neurolab
Visual System



SensoMotoric Instr. Eye
Tracking



Avotec audio



Nordic Neurolab Electrostatic Audio
System



MRI-compatible miniature video
camera

Hypothesis always a good thing to have – may be based on theory or functional neuroanatomy or previous neuroimaging experiments

Helpful if hypothesis suggests a specific neuroanatomical locus for change in neural activity or a network of regions?

Important to understand limitations of fMRI temporal and spatial resolution – useful to consult with FMRI staff regarding experimental design and possibilities

Understand limitations of fMRI environment and how stimulus can be presented to subject and behavior and performance monitored.

Write protocol and submit to Scientific Review Board (SRB) and get feedback.

Get IRB approval . Submit protocol to NMRF Protocol Committee for MRI safety/sanity check (<http://intranet.nmrf.nih.gov/>)

Get scanner time (assigned by DIRP to individual units, sections, labs, or programs)

Ensure all research staff are trained and ,if support is needed, technologist coverage arranged.

Contact Scientific and Statistical Computing Core Facility for assistance with experimental design and planning your analysis (<http://afni.nimh.nih.gov/sscc>)

Additional Resources for fMRI/Research

Functional MRI Facility

In-vivo NMR Facility

Scientific Statistical Computing Core

Neurophysiological Imaging Facility

MEG

Section on Instrumentation Core Facility

Thanks / Questions ?